PATENT

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

APPLICANTS: CHOI ET AL-1 (PCT)

PCT NO.: PCT/KR2005/002170

FILED: JULY 6, 2005

TITLE: DIAGNOSIS METHOD AND KITS FOR INHERITED NEUROPATHIES CAUSED BY DUPLICATION OR DELETION OF CHROMOSOME 17P11.2-P12 REGION

INFORMATION DISCLOSURE STATEMENT

MAIL STOP PCT
Commissioner of Patents
P.O. Box 1450
Alexandria, VA 22313-1450

Dear Sir:

Applicants are enclosing Form PTO-1449 disclosing the references cited in the International Search Report, copy enclosed. Copies of the U.S. references are not enclosed herewith, pursuant to current U.S.P.T.O. guidelines. The *Berger et al.* reference was discussed on page 2 of the Specification. The *Sereda, et al.* article, *Passage, et al.* article and the *Mersiyanova et al.* article were all discussed on page 5 of the Specification. The *Yoshihara et al.* article and the *Numakura et al.* article were discussed on page 6 of the Specification. Since the instant Information Disclosure Statement is being filed concurrently with the application, no official fee is required in connection with the same. It is respectfully requested that the foregoing Information Disclosure Statement be incorporated into the official file of the

concurrently-filed PCT patent application.

Respectfully submitted,

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Enclosures: PTO-1449 form, International Search Report and 2 references

ECR:lgh

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I hereby certify that this paper or fee is being deposited with the United States Postal Service "Express Mail Post Office to Addressee" service under 37 C.F.R. § 1.10, on the date indicated above, and is addressed to Commissioner of Patents, P.O. Box 1450, Alexandria, VA 22313-1450.

Kelly Espitia

FORM PTO-1449 (REV. 7-80)	U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE	ATTY. DOCKET NO.: CHOI ET AL--1 PCT	SERIAL NO: 10/589328
LIST OF REFERENCES CITED BY APPLICANT (Use several sheets if necessary)		APPLICANT: Byung Ok CHOI ET AL.	
		FILING DATE:	GROUP:

U.S. PATENT DOCUMENTS

*EXAMINER INITIAL		DOCUMENT NUMBER	DATE	NAME	CLASS	SUBCLASS	FILING DATE IF APPROPRIATE
	AA	5,780,223 A	07/1998	Lupski et al. (ISR)			
	AB	6,110,670 A	08/2000	Van Broeckhoven et al (ISR)			
	AC	5,306,616 A	04/1994	Lupski et al. (ISR)			
	AD	5,645,993 A	07/1997	Chance et al (ISR)			

OTHER REFERENCES *(Including Author, Title, Date, Pertinent Pages, Etc.)*

	AR		Vance JM, et al., "Localization of Charcot-Marie-Tooth disease type 1a (CMT1A) to chromosome 17p11.2', In: Genomics, Apr. 1991, Vol.9(4), pp. 623-628. (ISR) Enclosed
	AS		Kim SM, et al., "Hereditary neuropathy with liability to pressure palsies (HNPP) patients of Korean ancestry with chromosome 17p.11.2-p12 deletion', In: Exp Mol Med., 29 Feb. 2004, Vol. 36(1), pp. 28-35. (ISR) Enclosed
	AT		Berger P., Young P., Suter U. Neurogenet. 4: 1-15 (2002). Spec-to follow
	AU		Sereda M.W., Horste G.M., Suter U., Uzma N., Nave K.-A. Nature Genet. 9: 1533-1537 (2003) Spec-to follow
	AV		Passage E., Norreel J.C., Noach-Fraissignes P., Sanguedolce V., Pizant J., Thirion X., Robaglia-Schlupp A., Pellissier J.F., Fontes M. Nature Genet. 10: 396-401 (2004) Spec-to follow
	AW		Mersiyanova I.V., Ismailov S.M., Polyakov A.V., Dadali E.L., Fedotov V.P., Nelis E., et al. Human Mutat. 15: 340-347 (2000) Spec-to follow
	AX		Yoshihara T., Yamamoto M., Doyu M., Misu K.I., Hattori N. Hasegawa Y., Mokuno K., Mitsuma T., Sobue G. Hum. Mutat. 16: 177-178 (2000) Spec-to follow
	AY		Numakura C., Lin C., Ikegami T., Guldberg P., Hayasaka K Human Mutat. 20: 392-398 (2002) Spec- to follow

EXAMINER	DATE CONSIDERED

*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609; draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.